

んづう

PATENT Customer No. 22,852 Attorney Docket No. **6832.0013**

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of:	
Craig A. ROSEN et al.) Group Art Unit: 1653
Application No.: 09/832,929)) Examiner: Hope A. Robinson
Filed: April 12, 2001	RECEIVED
For: ALBUMIN FUSION PROTEINS)
Commissioner for Patents	AUG 1 8 2003
P.O. Box 1450 Alexandria, VA 22313-1450	TECH CENTER 1600/2900
Sir:	

RESPONSE TO RESTRICTION REQUIREMENT

In a restriction requirement dated August 4, 2003, the Examiner required restriction under 35 U.S.C. § 121 between:

Group I Claims 1-21, drawn to an albumin fusion protein comprising a therapeutic protein X and albumin (SEQ ID NO: 18);

Group II Claims 22-25, drawn to a method of treating a disease or disorder in a patient;

Group III Claim 26, drawn to a method of extending the shelf life of therapeutic protein X; and

Group IV Claims 27-29, drawn to a nucleic acid molecule.

FINNEGAN HENDERSON FARABOW GARRETT & DUNNER

1300 I Street, NW Washington, DC 20005 202.408.4000 Fax 202.408.4400 www.finnegan.com

Attorney Docket No.: 6832.0013

Application Serial No.: 09/832,929

Applicants provisionally elect to prosecute Group I, claims 1-21 drawn to an albumin fusion protein comprising a therapeutic protein X and albumin (SEQ ID NO: 18).

According to MPEP § 803, there are two requirements that must be met before a proper restriction requirement may be made: (1) the inventions must be independent or distinct as claimed; and (2) there must be a serious burden on the Examiner if restriction is not required. Applicants respectfully submit that the Office has failed to establish the second requirement set forth in MPEP § 803, that a serious burden exists on the Examiner if restriction is not required between the Groups of claims.

In the present invention, Group I is directed to an albumin fusion protein comprising a therapeutic protein X and albumin. Group II is directed to a method of treating a disease or disorder comprising the step of administering the albumin fusion protein as defined in Group I. Additionally, Group IV is directed to a nucleic acid molecule encoding the albumin fusion protein as defined in Group I. A search and examination of the subject matter of Group I would encompass a search for the subject matters of Groups II and IV, and any additional search would not impose a serious burden upon the Examiner.

It is therefore respectfully requested that the restriction requirement be reconsidered. In the event that the restriction requirement is maintained, Applicants reserve the right to file divisional applications on the non-elected inventions and/or to request rejoinder of appropriate claims once the subject matter of claims 1-21 is found allowable.

FINNEGAN HENDERSON FARABOW GARRETT & DUNNER LLP

1300 I Street, NW Washington, DC 20005 202.408.4000 Fax 202.408.4400 www.finnegan.com

Attorney Docket No.: 6832.0013 Application Serial No.: 09/832,929

Please grant any extensions of time required to enter this response and charge any additional required fees to our deposit account 06-0916.

Respectfully submitted,

FINNEGAN, HENDERSON, FARABOW, GARRETT & DUNNER, L.L.P.

By: Charles E. Van Horn
Reg. No. 40,266

Dated: August 15, 2003

FINNEGAN HENDERSON FARABOW GARRETT & DUNNER LLP

1300 I Street, NW Washington, DC 20005 202.408.4000 Fax 202.408.4400 www.finnegan.com